Serial No.: New - PCT/ JP2004/0017779 Nat'l Phase

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The following Listing of Claims will replace all prior versions, and listings, of claims

in the application.

**LISTING OF CLAIMS:** 

1. (Original) A method of assaying whether a chemical is present in a test

specimen or not, comprising culturing a gene-disrupted strain of a microorganism in the

presence of the test specimen, and using cell response of the gene-disrupted strain to the

chemical as an index.

2. (Original) The method according to claim 1, wherein

the cell response of a gene-disrupted strain to a chemical is life or death of a cell,

and/or proliferation ability, aspiration amount, enzyme activity and/or a change in gene

expression.

3. (Original) The method according to claim 2, wherein

the change in gene expression is a change in a RNA amount or a mRNA amount.

4. (Original) The method according to claim 2, wherein

the change in gene expression is measured by reporter gene assay.

5. (Currently Amended) The method according to any one of claims 1 to 4 claim

1, wherein

the microorganism is yeast.

Page 9 of 15

Serial No.: New – PCT/ JP2004/0017779 Nat'l Phase

Filed: Herewith

6. (Currently Amended) The method according to claim 5, wherein a gene to be disrupted is classified into:

amino acid metabolism (01.01), nitrogen and sulfur metabolism (01.02), nucleotide metabolism (01.03), phosphate metabolism (01.04), C-compound and carbohydrate metabolism (01.05), lipid, fatty acid and isoprenoid metabolism (01.06), metabolism of vitamins, cofactors and prosthetic groups (01.07) of metabolism (01);

DNA processing (03.01), cell cycle (03.03) of cell cycle and DNA processing (03); mRNA transcription (04.05), RNA transport (04.07) of transcription (04); ribosome biosynthesis (05.01), translation control (05.07) of protein synthesis (05); protein targeting, sorting, translocation (06.04), protein modification (06.07), assembly of protein complex (06.10), proteolysis (06.13) of protein fate (06);

nuclear transport (08.01), vesicular transport (Golgi network etc) (08.07), vacuolar transport (08.13), cellular import (08.19), cytoskeleton-dependent transport (08.22), other intracellular transport activities (08.99) of intracellular transport and transport mechanism (08);

stress response (11.01), detoxification (11.07) of cell rescue, defense and pathogemicity (11);

ionic homeostasis (13.01), cell sensitivity and response (13.11) of intracellular environmental regulation/interaction (13);

cell growth/morphogenesis (14.01), cell differentiation (14.04) of cell fate (14); cell wall (30.01), cytoskeleton (30.04), nucleus (30.10), mitochondria (30.16) of cell tissue control (30);

Serial No.: New - PCT/ JP2004/0017779 Nat'l Phase

Filed: Herewith

ion transporter (67.04), vitamin/cofactor transporter (67.21), transport mechanism (67.50), other transport promotion (67.99) of transport promotion (67); unclassified (98); and/or unclassified protein (99).

- 7. (Original) The method according to claims 6, wherein the gene to be disrupted is involved in a vacuole.
- 8. (Currently Amended) The method according to claim 6, wherein the metabolism (01) gene to be disrupted is YGL026C, YGR180C, YDR127W, YCR028C, YLR284C, YOR221C, YAL021C, YGL224C, YBL042C, YDR148C, YHL025W, YLR307W, YLR345W, YLR354C, YPL129W, or YPR060C.
- (Currently Amended) The method according to claim 6, wherein the cell cycle and DNA processing (03) gene to be disrupted is YGR180C,
  YDR150W, YGL240W, YBL058W, YIL036W, YLR226W, YLR381W, YOR026W,
  YPL018W, YBL063W, YDR363W-A, YIR026C, YLR234W, YMR032W or YPL129W.
- 10. (Currently Amended) The method according to claim 6, wherein the transcription (04) gene to be disrupted is YGR006W, YIL036W, YKR082W, YLR226W, YML112W, YMR021C, YAL021C, YDR195W, YOL068C, YBR279W, YGL070C, YGL071W, YGL222C, YHL025W, YLR266C or YPL129W.

Serial No.: New - PCT/ JP2004/0017779 Nat'l Phase

Filed: Herewith

11. (Currently Amended) The method according to claim 6, wherein the protein synthesis (05) gene to be disrupted is YBL058W, YLR287C-A, YGR084C or YLR344W.

- 12. (Currently Amended) The method according to claim 6, wherein the protein fate (06) gene to be disrupted is YKL080W, YLR447C, YGL240W, YGR105W, YGL206C, YKL119C, YDR414C, YHR060W, YLR292C, YLR306W, YGL227W or YGR270W.
- 13. (Currently Amended) The method according to claim 6, wherein the intracellular transport and transport mechanism (08) gene to be disrupted is YPR036W, YDR027C, YHR039C, YKL080W, YLR447C, YGL206C, YKR082W, YLR292C or YBL063W.
  - 14. (Currently Amended) The method according to claim 6, wherein the detoxification (11) gene to be disrupted is YJR104C or YMR021C.
- 15. (Currently Amended) The method according to claim 6, wherein the intracellular regulation/interaction (13) gene to be disrupted is YPR036W, YHR039C-B, YKL080W, YLR447C, YGL071W or YIR026C.
  - 16. (Currently Amended) The method according to claim 6, wherein

Serial No.: New – PCT/ JP2004/0017779 Nat'l Phase

Filed: Herewith

the cell fate (14) gene to be disrupted is YDL151C, YBL058W, YKR082W, YDL151C, YOL068C, YDR363W-A, YHL025W, YIR026C, YLR307W, YMR032W or YPL129W.

- 17. (Currently Amended) The method according to claim 6, wherein the cell tissue control (30) gene to be disrupted is YDR027C, YDR414C, YLR381W, YGR084C or YMR032W.
- 18. (Currently Amended) The method according to claim 6, wherein the transport promotion (67) gene to be disrupted is YPR036W, YHR026W, YHR039C, YKL080W, YLR447C, YCR028C or YLR292C.
  - 19. (Currently Amended) The method according to claim 6, wherein the unclassified (98) gene to be disrupted is YBL056W.
- 20. (Currently Amended) The method according to claim 6, wherein the unclassified protein (99) gene to be disrupted is YDR149C, YLR285W, YLR311C, YOR331C, YPR123C, YDR525W-A, YDR539W, YDR540C, YGL246C, YJL204C, YLR282C, YLR287C, YLR290C, YJL188C, YJL192C, YJL211C, YKL037W, YLR283W, YLR312C, YLR315W, YLR320W or YPL030W.
- 21. (Original) A kit containing a gene-disrupted strain of a microorganism, which is used for detecting whether a chemical is present in a test specimen or not.

Serial No.: New – PCT/ JP2004/0017779 Nat'l Phase

Filed: Herewith

22. (Original) A composition containing a gene-disrupted strain of a microorganism, for detecting whether a chemical is present in a test specimen or not.

23. (Original) Use of a gene-disrupted strain of a microorganism, for detecting whether a chemical is present in a test specimen or not.